

# SAHA Patient: IV vs ORAL (200 mg/dose)

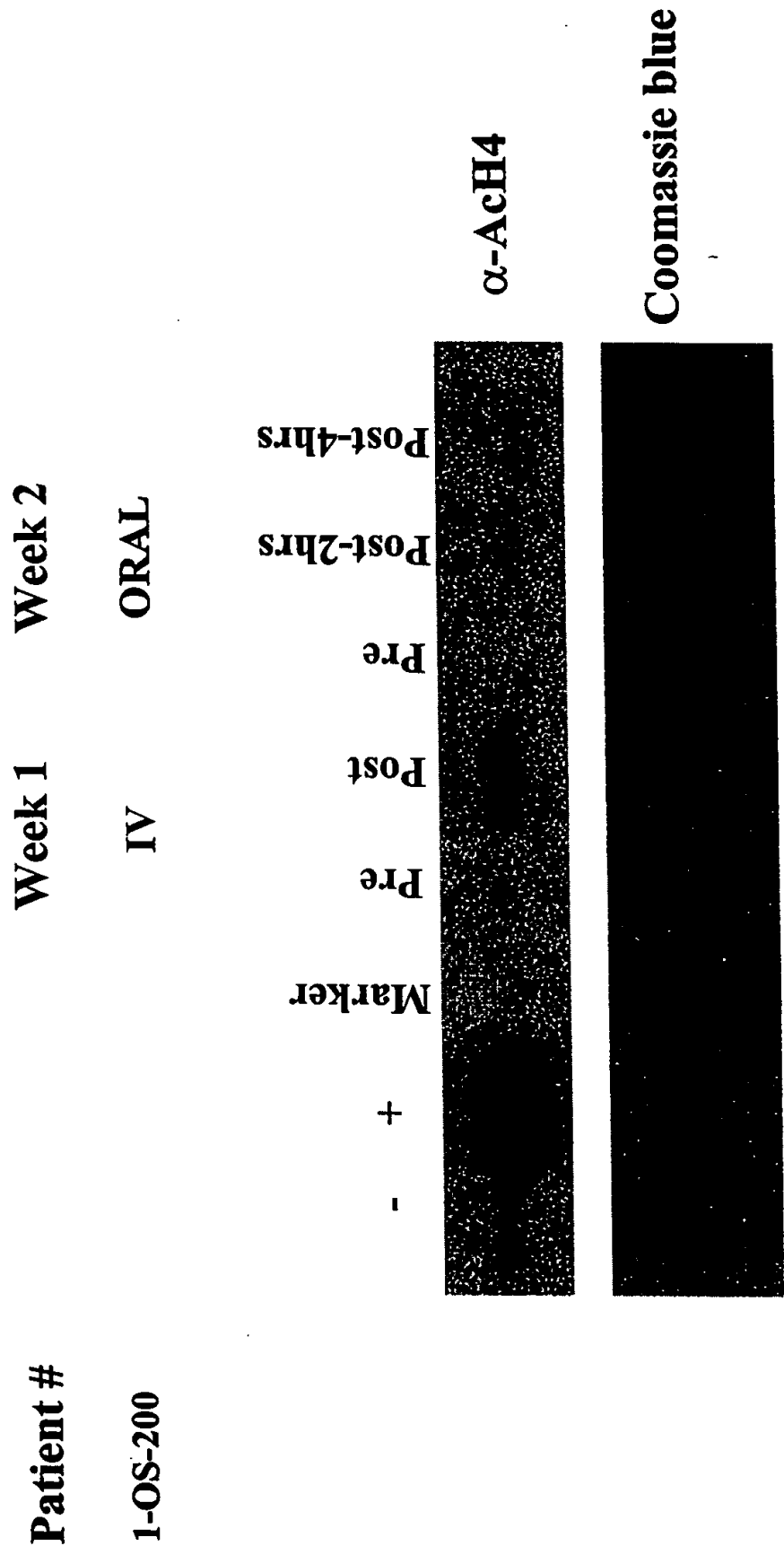


FIG. 1

**Protocol 01-021 (ORAL SAHA) ARM A: SOLID  
TUMOR PATIENTS Cohort I (200 mg/dose)**

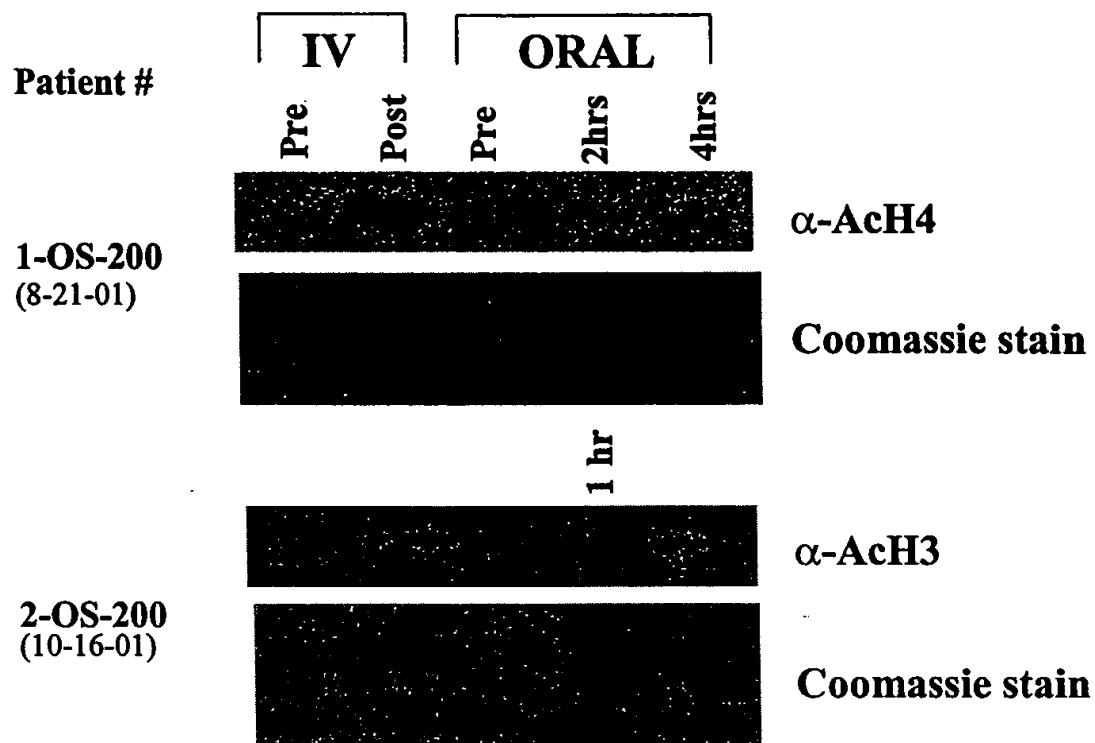


FIG. 2

# SAHA Patients: ORAL /Cohort I (200 mg/dose)

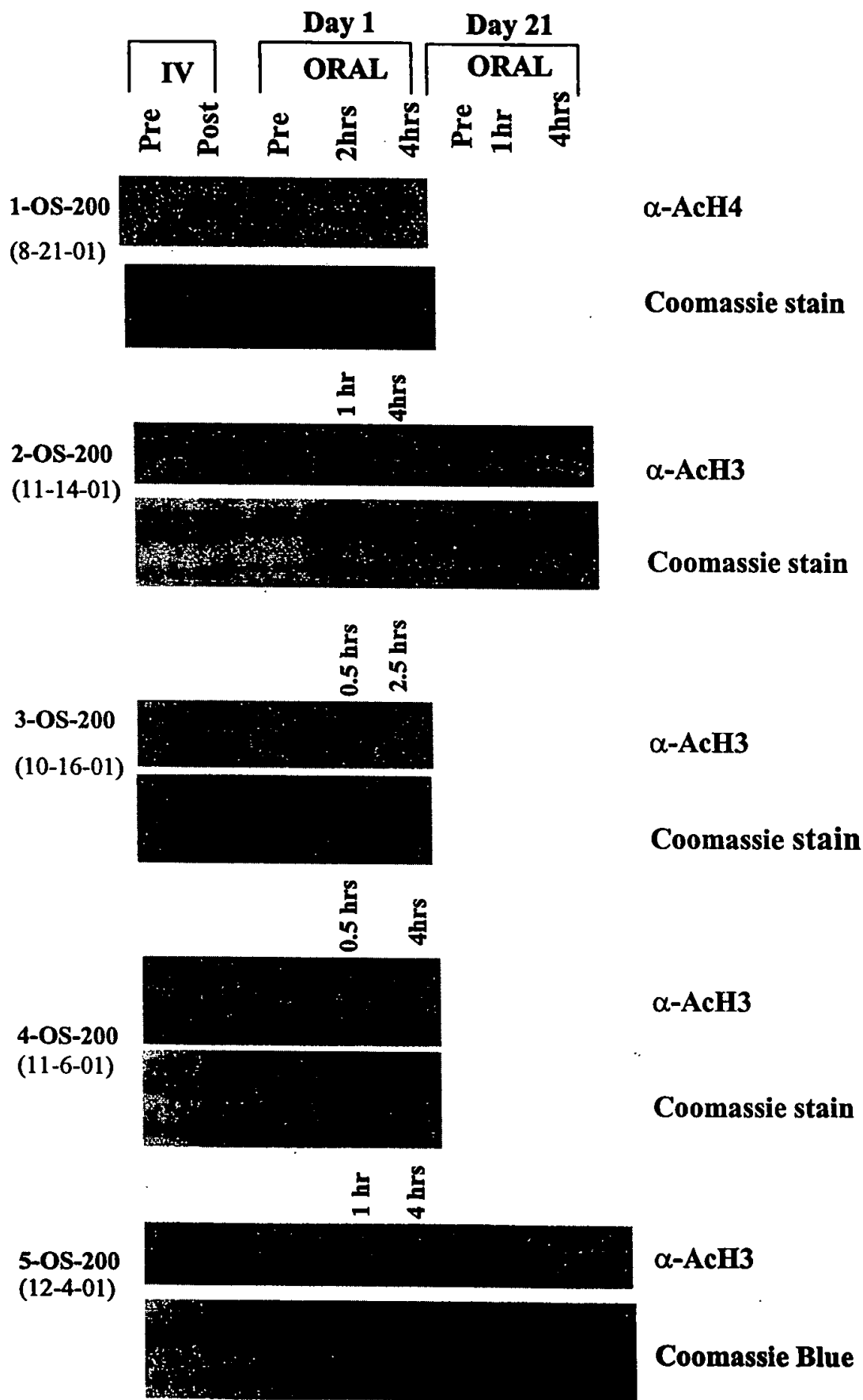


FIG. 3

**Protocol 01-021 (ORAL SAHA)**  
**ARM A: SOLID TUMOR PATIENTS**  
**Cohort I (200 mg/dose)**

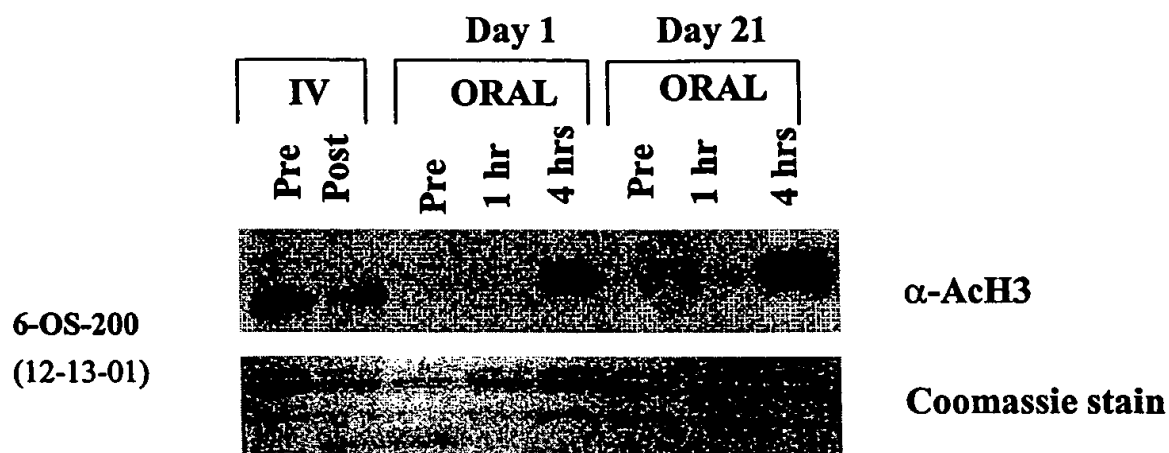
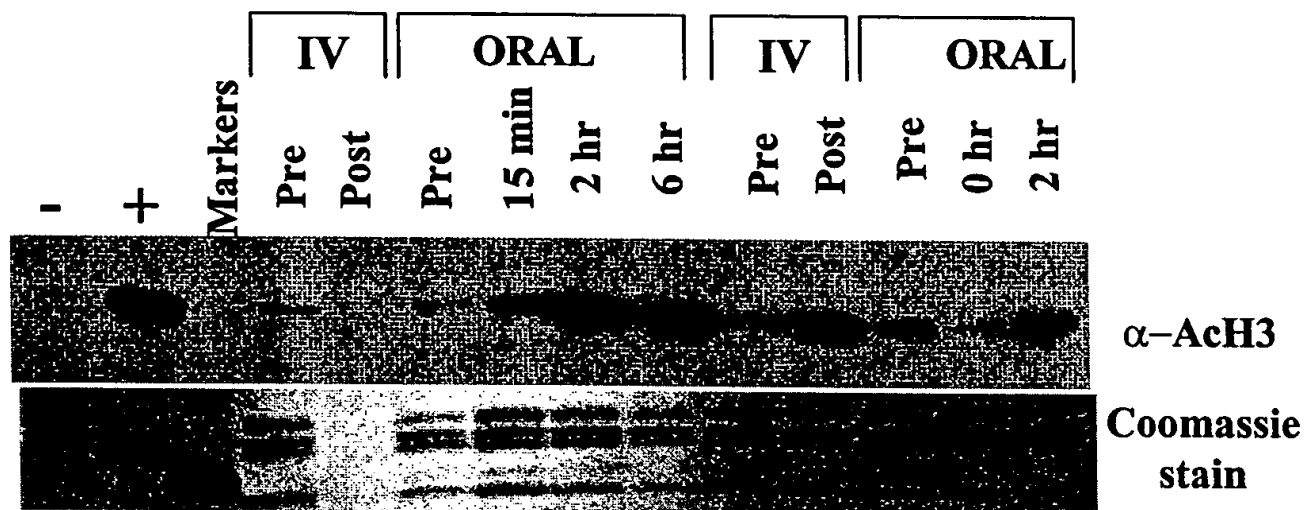


FIG. 4

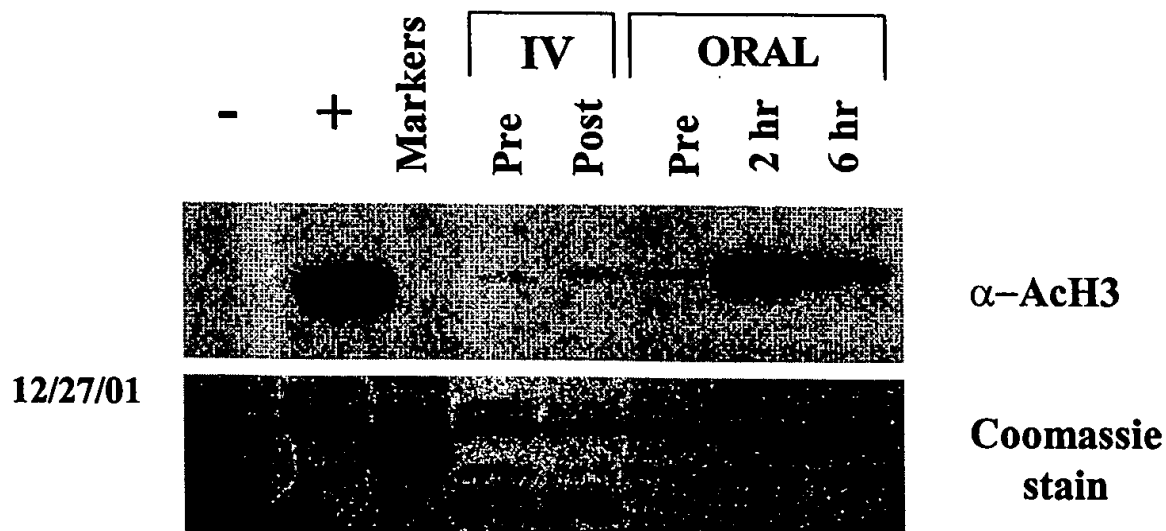
**Protocol 01-021 (ORAL SAHA)**  
**ARM A: SOLID TUMOR PATIENTS**

**Cohort IIa (400 mg/dose)**  
**7-OS-400                      8-OS-400**



12/19/01

**9-OS-400**



12/27/01

**FIG. 5**

# SAHA Patients: ORAL/Cohort II (400 mg/dose)

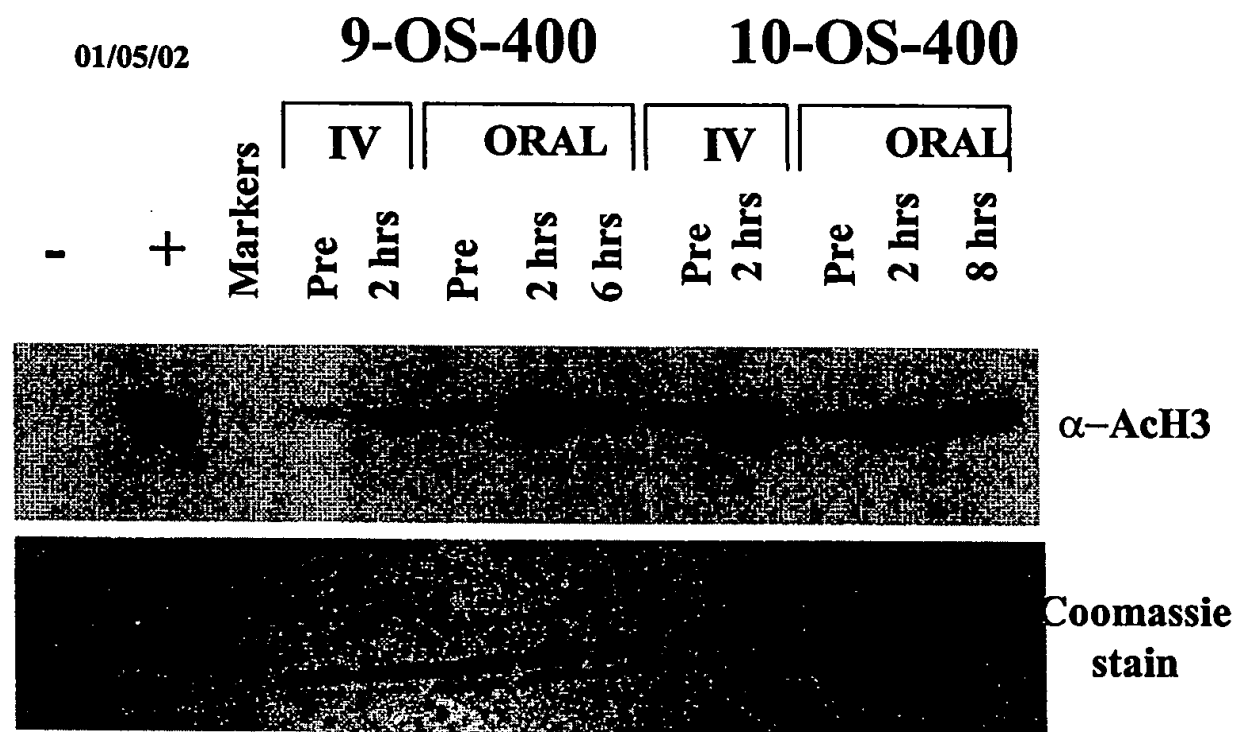
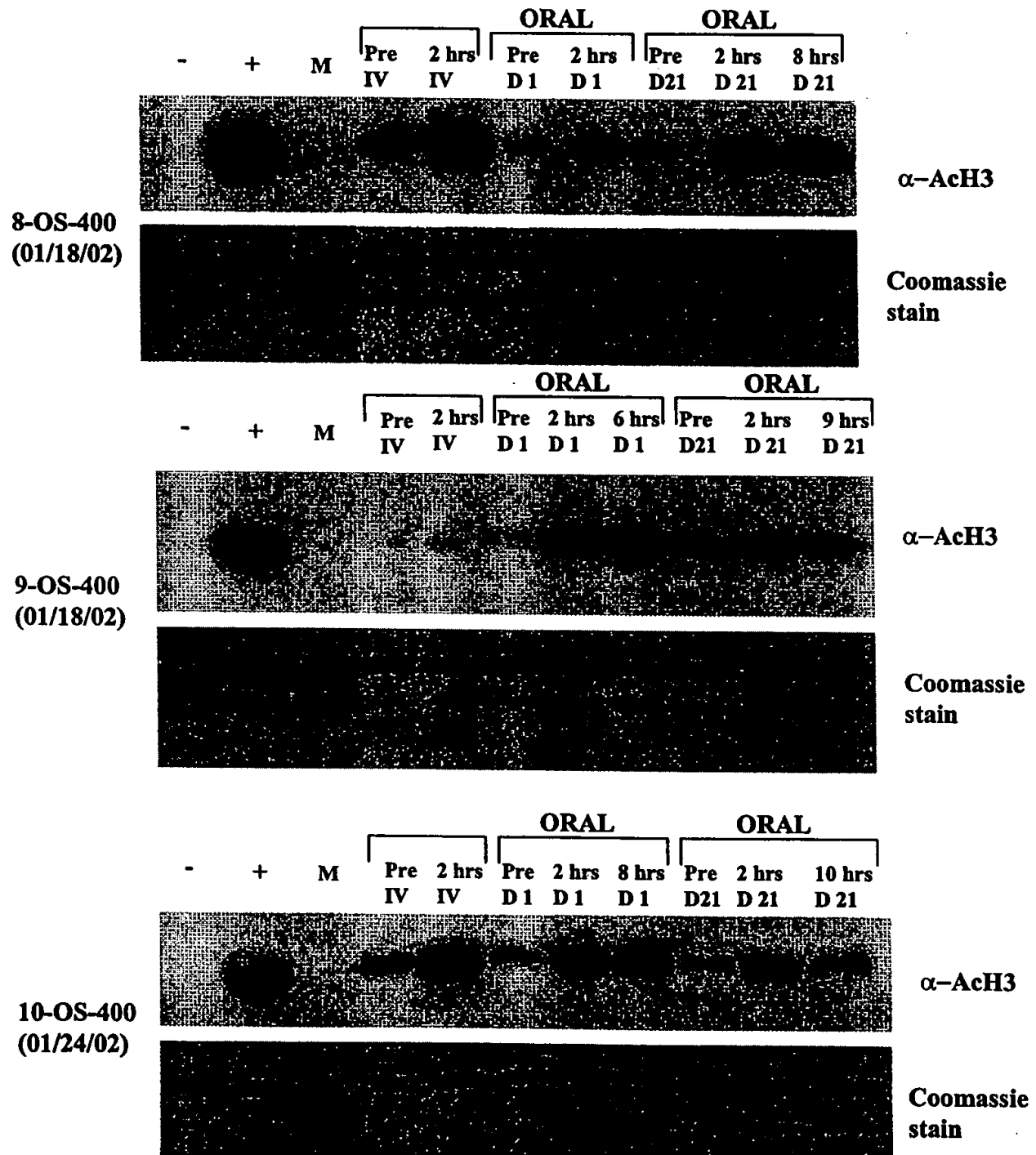


FIG. 6

**Protocol 01-021 (ORAL SAHA )**  
**ARM A: SOLID TUMOR PATIENTS**  
**Cohort IIa (400 mg/dose)**



**FIG. 7**

# SAHA Patients: ORAL/Cohort II (400 mg/dose)

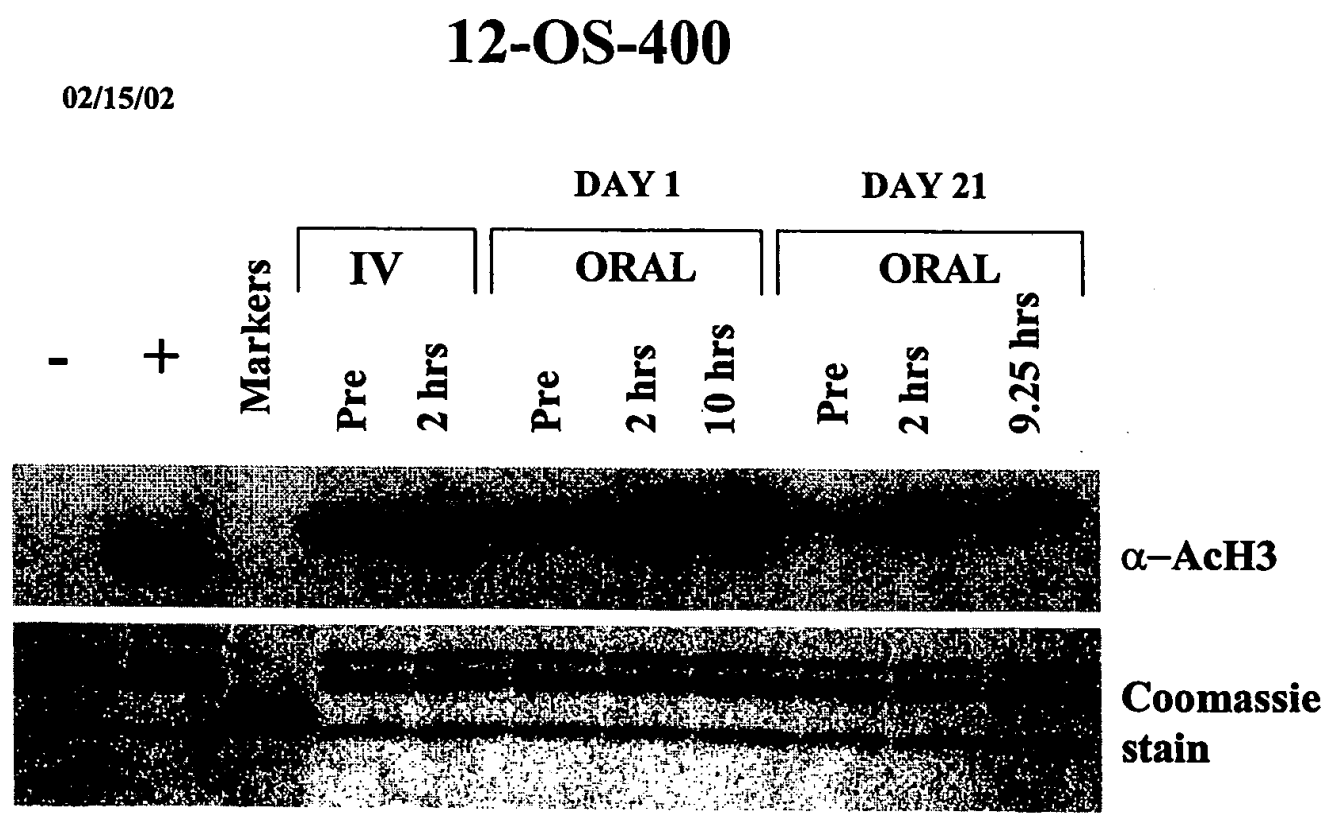
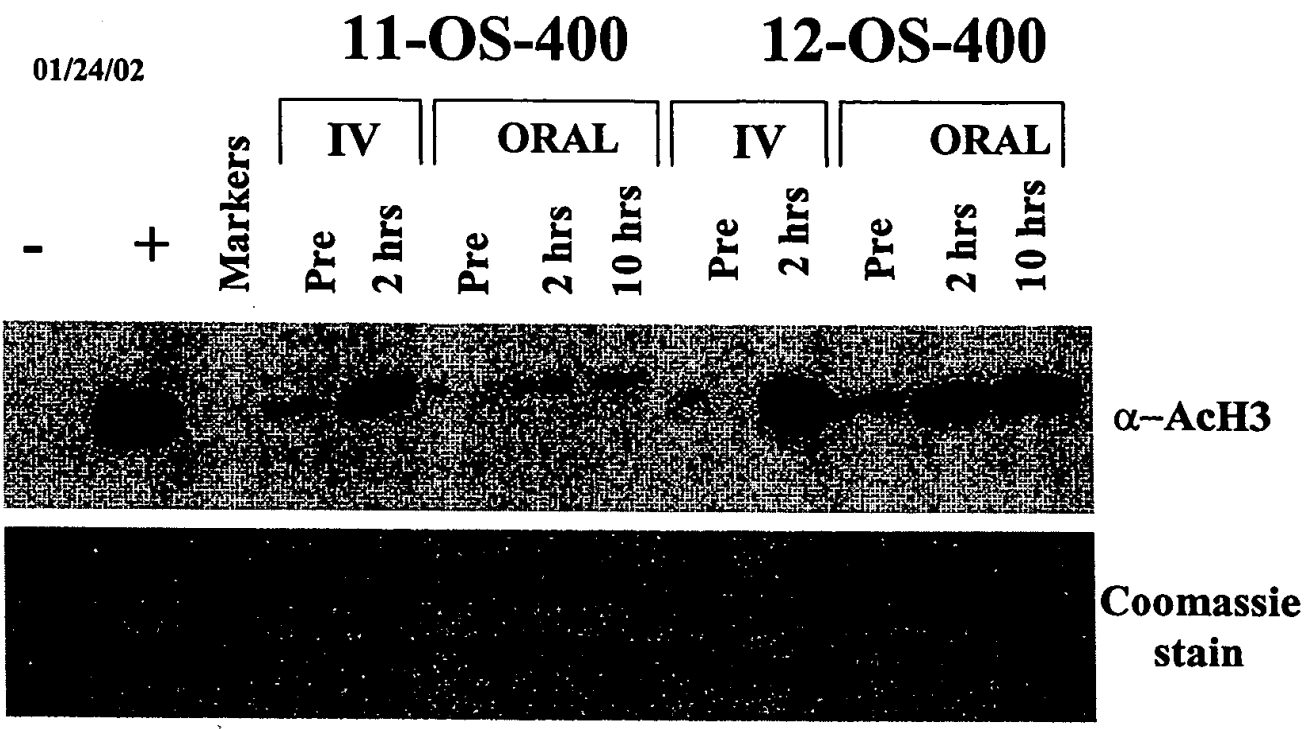


FIG. 8



# Oral 200 mg vs. 400 mg (Fasting)

Oral dose on Cycle 1 Day 8

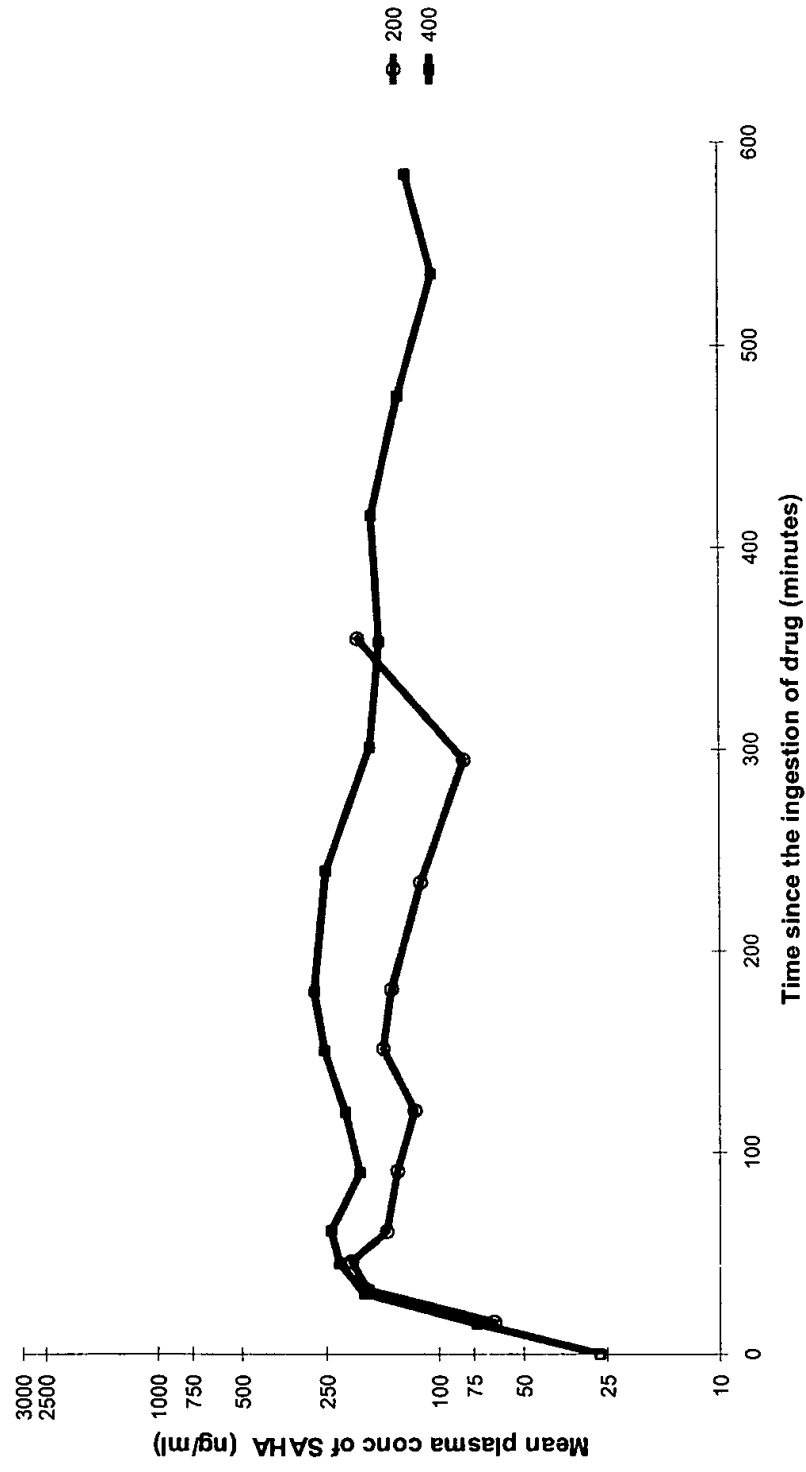


Figure 9A

# Oral 200 mg vs. 400 mg (no-fasting)

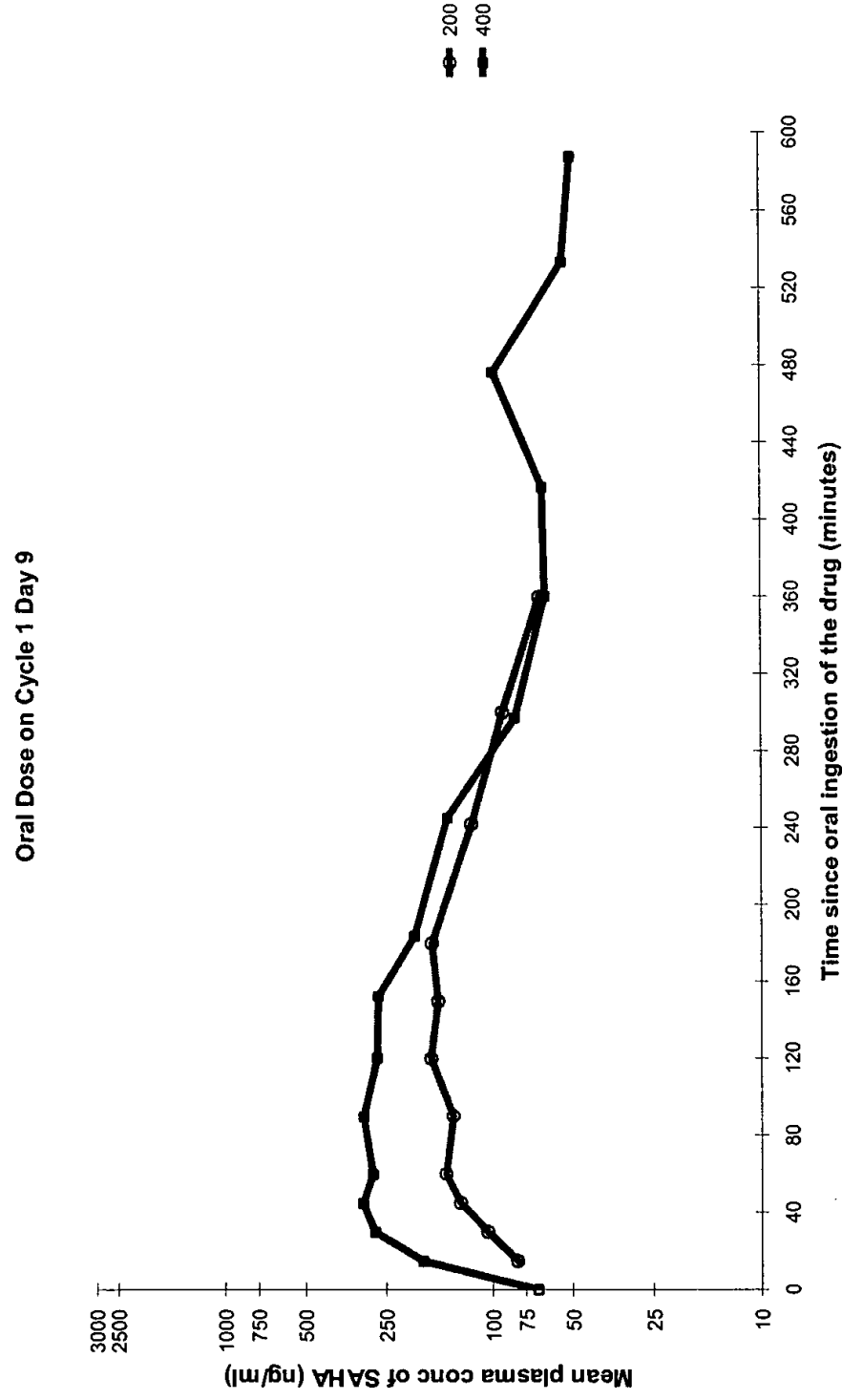


Figure 9B

# IV 200 mg vs. 400 mg

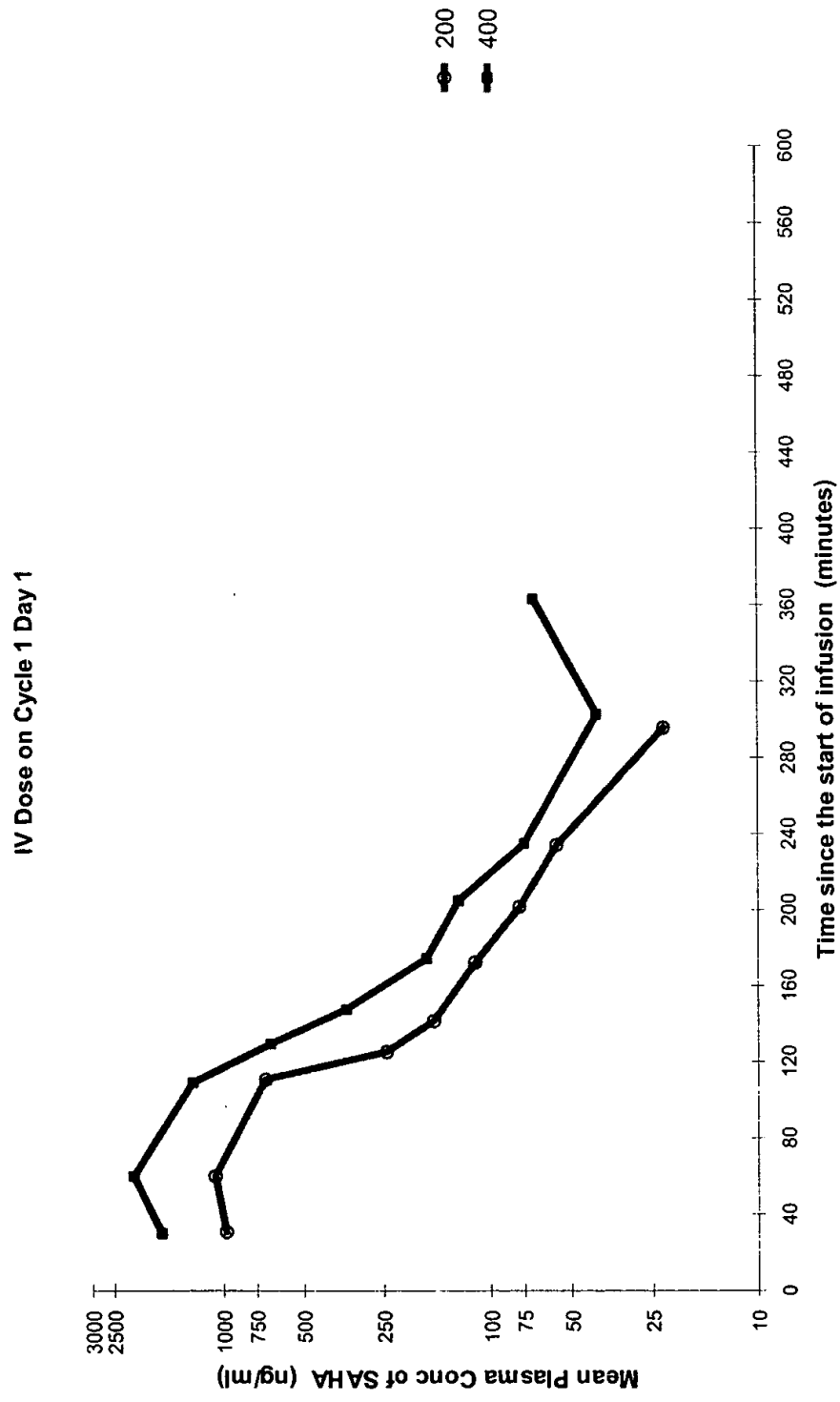
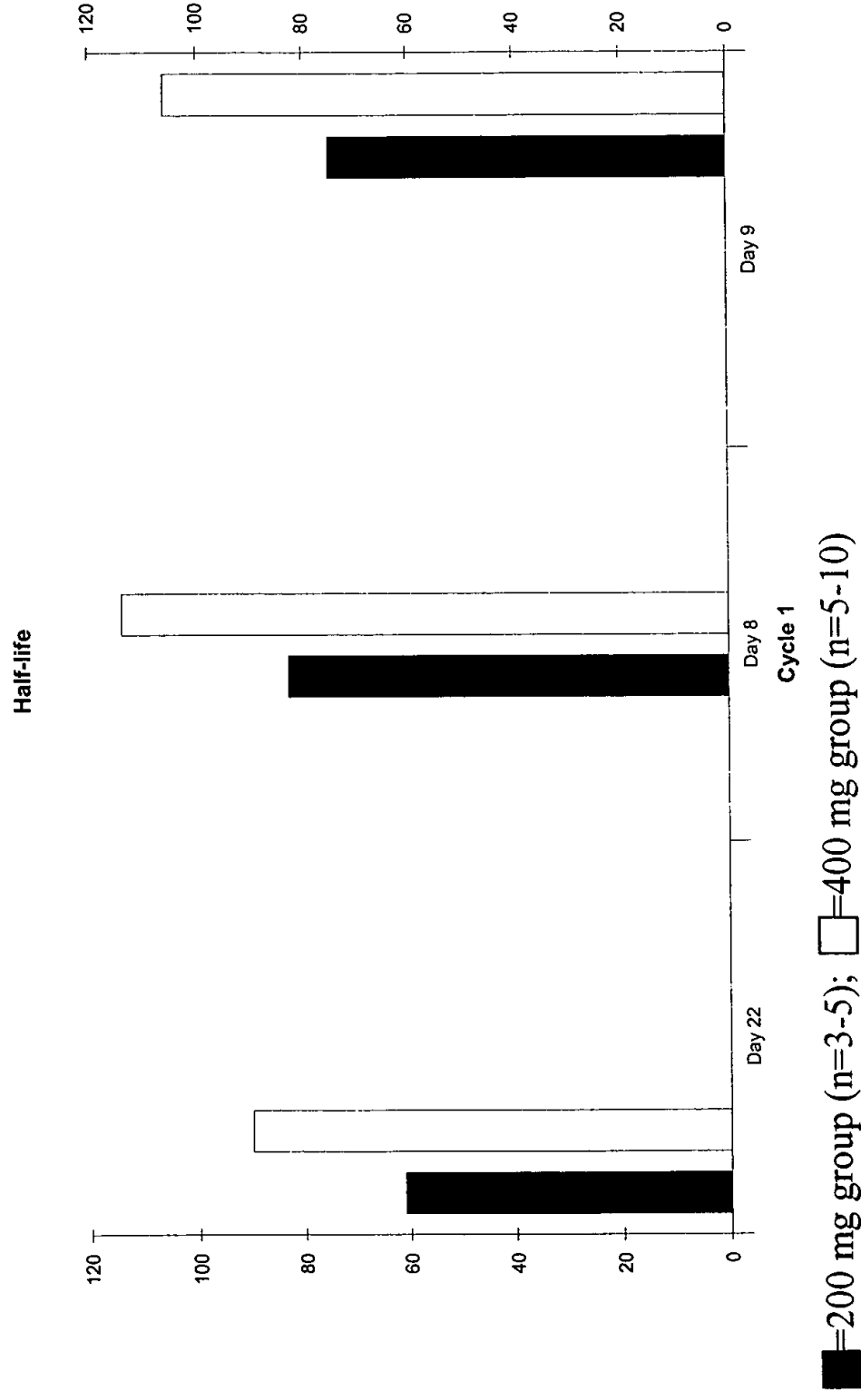


Figure 9C

# Apparent Half-life of the Oral Dose

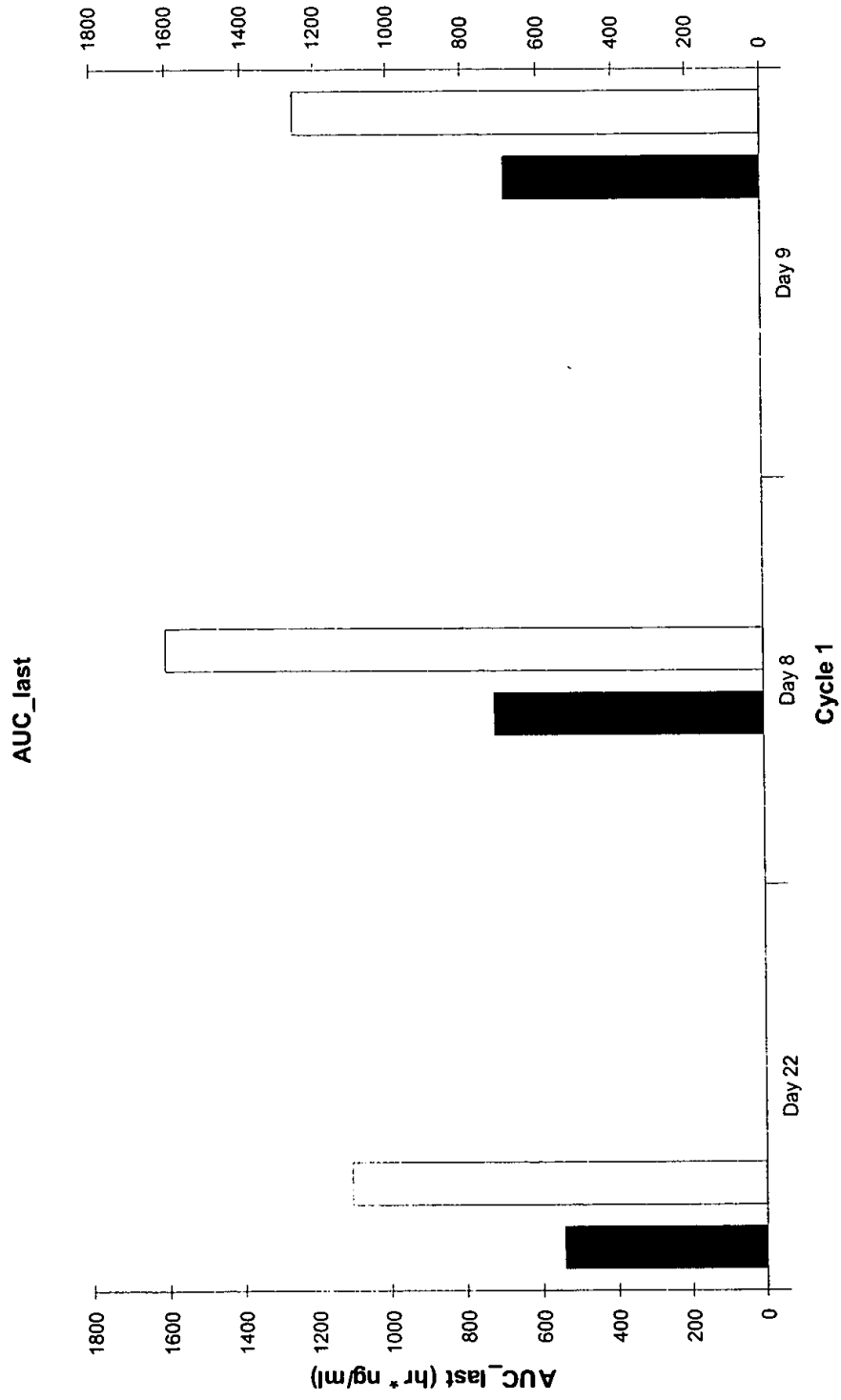
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Figure 10

# AUC of the Oral Dose



■ = 200 mg group (n=3-5); □ = 400 mg group (n=5-10)

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Figure 11

# Bioavailability

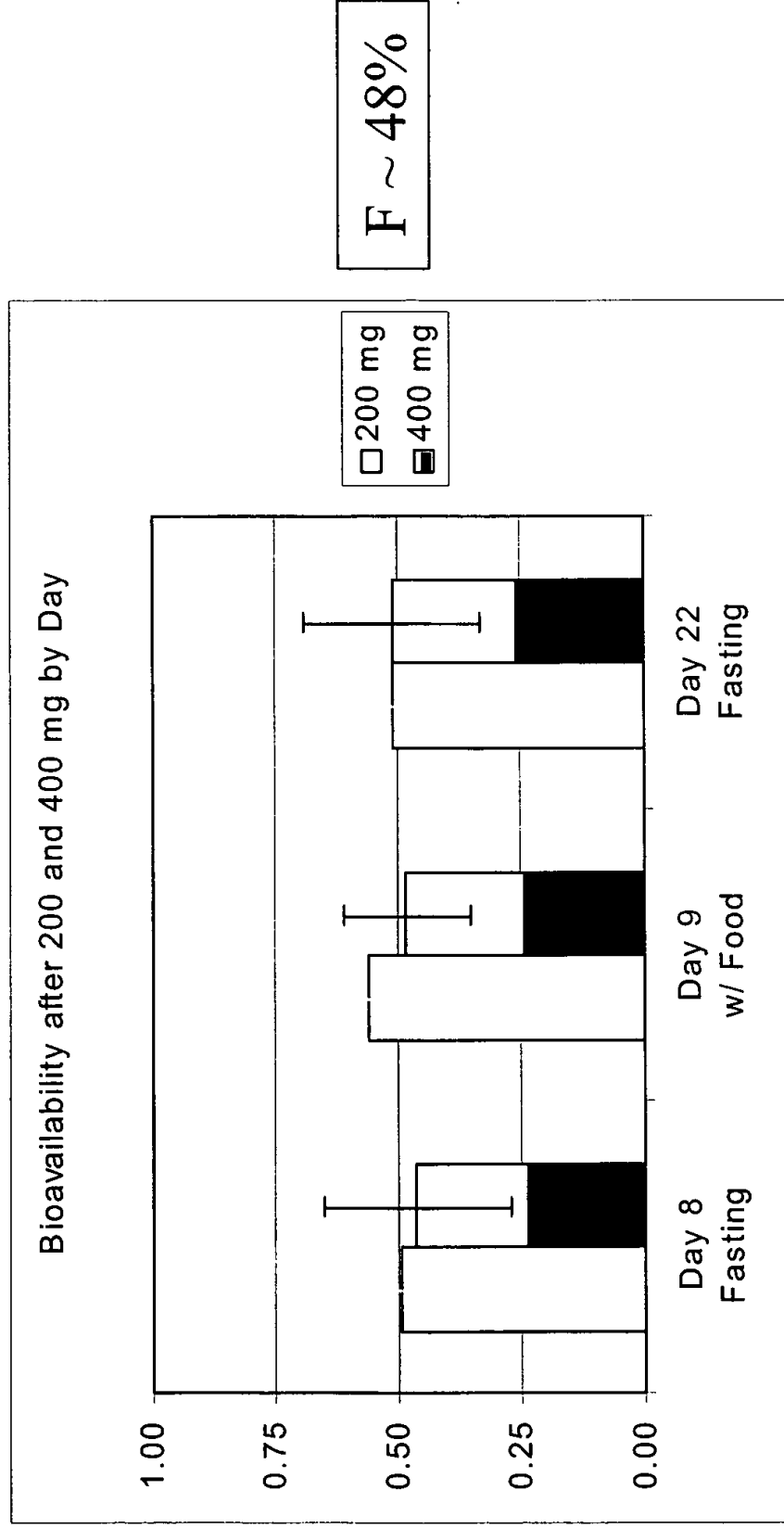


Figure 12